

DEPARTMENT OF HEALTH AND HUMAN SERVICES  
FOOD AND DRUG ADMINISTRATION

INFUSION PUMP INFORMATION SUBMISSION REPORT

See Burden Statement on back of form.  
Form Approved: OMB No. 0910-0387, Expiration Date: March 31, 2005.

MANUFACTURER NAME	PUMP BRAND NAME	PUMP MODEL NUMBER	FDA SUBMISSION NO. <i>(for FDA)</i>	YEAR MARKETED <i>(for FDA)</i>	PREVIOUS MANUFACTURER(S)
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**INFUSION PUMP INFORMATION SUBMISSION REPORT** *(continued)*

CONTACT PERSON <i>(for FDA)</i>	CONTACT PHONE NO. <i>(for FDA)</i>	CONTACT PHONE NO. <i>(for Consumers)</i>	MAILING ADDRESS	CITY	STATE	ZIP CODE	WEBSITE ADDRESS
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**Public reporting burden** Public reporting burden for this collection of information is estimated to average 1-2 hours per submission form, including the time for reviewing instructions, searching existing data sources, adhering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to:

Department of Health and Human Services  
Food and Drug Administration  
Center for Devices and Radiological Health  
1350 Piccard Drive, 340N  
Rockville, MD 20850